Appl. No. 10/088,970 Amdt. dated January 18, 2006 Reply to Final Office Action of Dec. 21, 2005

PATENT

Amendments to the Claims:

This listing of claims will replace all prior versions, and listings, of claims in the application:

Listing of Claims:

- 1. (currently amended) A method for <u>diagnosing prostate cancer versus benign prostate</u>
 <u>hyperplasia</u> aiding a prostate cancer diagnosis, the method comprising:
- i. obtaining from a subject a sample containing a plurality of prostrate related protein markers having apparent molecular weights below 10,000 Da;
- ii. determining by mass spectroscopy a test amount of the plurality of protein markers in the sample, the protein markers having an apparent molecular weight of less than 10,000 Da;
- iii. comparing the test amount of the plurality of protein markers having apparent molecular weight of less than 10,000 Da with an amount of a plurality of protein markers having an apparent molecular weight of less than 10,000 Da from a control sample where the control sample originates from benign prostate hyperplasia; and
- iv. determining whether the test amount is a diagnostic amount consistent with a diagnosis of prostate cancer versus benign prostate hyperplasia.

Claims 2-7 (canceled)

8. (original) The method of claim 1, wherein the sample is selected from the group consisting of blood, serum, urine, semen, seminal fluid, seminal plasma, and tissue extracts.

Claims 9-11 (canceled)

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- 12. (previously presented) The method of claim 1, the method further comprising:
- (a) generating data on the sample with the mass spectrometer indicating intensity of signal for mass/charge ratios;
 - (b) transforming the data into computer-readable form; and
- (c) operating a computer to execute an algorithm, wherein the algorithm determines closeness-of-fit between the computer-readable data and data indicating a diagnosis of prostate cancer or a negative diagnosis.

Claims 13-19 (canceled)

20. (previously presented) The method of claim 1, wherein the sample is seminal plasma.

Claims 21-83 (canceled)

- 84. (previously presented) The method of claim 1 where the protein markers are adsorbed onto a probe comprising an adsorbent of a hydrophilic polymer.
- 85. (previously presented) The method of claim 1 where the protein markers are adsorbed onto a probe comprising a metal binding group.
- 86. (previously presented) The method of claim 84 where the adsorbent comprises a hydrophobic group.
- 87. (previously presented) The method of claim 84 where the adsorbent comprises a cationic group.
- 88. (previously presented) The method of claim 84 where the adsorbent comprises a metal ion chelating group.

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- 89. (new) The method of claim 20, the method further comprising:
- (a) generating data on the sample with the mass spectrometer indicating intensity of signal for mass/charge ratios;
 - (b) transforming the data into computer-readable form; and
- (c) operating a computer to execute an algorithm, wherein the algorithm determines closeness-of-fit between the computer-readable data and data indicating a diagnosis of prostate cancer or a negative diagnosis.
- 90. (new) The method of claim 20 where the protein markers are adsorbed onto a probe comprising an adsorbent of a hydrophilic polymer.
 - 91. (new) The method of claim 20 where the protein markers are adsorbed onto a probe comprising a metal binding group.
 - 92. (new) The method of claim 90 where the adsorbent comprises a hydrophobic group.
 - 93. (new) The method of claim 90 where the adsorbent comprises a cationic group.
- 94. (new) The method of claim 90 where the adsorbent comprises a metal ion chelating group.